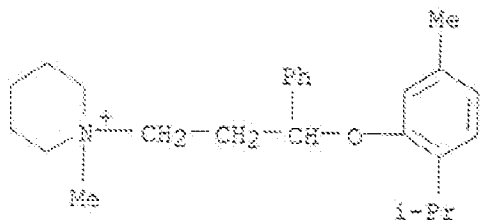


AN 1977:189458 CAPLUS
 DN 86:189458
 TI Aromatic amino ether quaternary ammonium salts
 IN Ogawa, Shuntaro; Morita, Kan; Yoshida, Akiyoshi
 PA Rokko Pharmaceutical Co., Ltd., Japan
 SO Japan., 9 pp.
 CODEN: JAXXAD
 DT Patent
 LA Japanese
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
FI	JP 51044934	B4	19761201	JP 1969-99283	19691209
AB	<p>RC6H4ZCHPhCH2CH2N+R1R2R3 X- (I; R = H, halo, Me; R1, R2 = H, alkyl, R1R2 = alkylene contg. optional O atom; R3 = alkyl; X = halo; Z = O, S) were prepd. by quaternization of RC6H4ZCHPhCH2CH2NR1R2 (II) with R3X. I were useful as antispasmodics, anticholinergics, antiinflammants, and analgesics. Thus, excess MeI was added to II [R = H, R1R2 = (CH2)5, Z = O], obtained from 2.4 g of its HCl salt after treatment with aq. NaOH and Et2O extn., in MeOH at room temp. to give 2.6 g I [R = H, R1R2 = (CH2)5, R3 = Me, X = iodo, Z = O], which had anticholinergic activity with ED50 of 1.6 .times. 10-6 g/mL in guinea pigs. Similarly prepd. were 17 addnl. I and their biol. activity given.</p>				
IT	42063-78-5P	<p>RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)</p>			
RN	42063-78-5	CAPLUS			
CN	<p>Piperidinium, 1-methyl-1-[3-[5-methyl-2-(1-methylethyl)phenoxy]-3-phenylpropyl]-, iodide (9CI) (CA INDEX NAME)</p>				

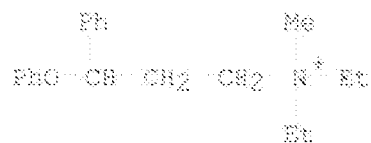


● I⁻

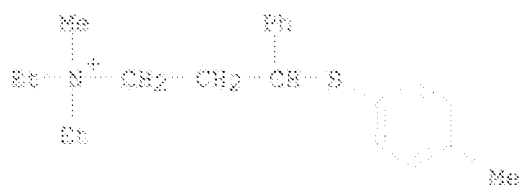
AN 1977:189458 CAPLUS
 DN 86:189458
 TI Aromatic amino ether quaternary ammonium salts
 IN Ogawa, Shuntaro; Morita, Ken; Yoshida, Akayoshi
 PA Robto Pharmaceutical Co., Ltd., Japan
 SO Japan., 9 pp.
 CODEN: JAXXAD
 OT Patent
 LA Japanese
 IC C07C093-12
 CC 25-4 (Noncondensed Aromatic Compounds)
 Section cross-reference(s): 27

FAN.CNT 1

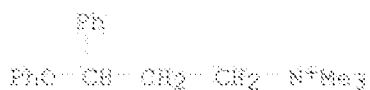
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	JP 51044934	B4	19761201	JP 1969-99283	19691209
AB	<p>RC6H4ZCHPhCH2CH2N+R1R2R3 X- (I; R = H, halo, Me; R1, R2 = H, alkyl, R1R2 = alkylene contg. optional O atom; R3 = alkyl; X = halo; Z = O, S) were prepd. by quaternization of RC6H4ZCHPhCH2CH2NR1R2 (II) with R3X. I were useful as antispasmodics, anticholinergics, antiinflammatories, and analgesics. Thus, excess MeI was added to II (R = H, R1R2 = (CH2)5, Z = O), obtained from 2.4 g of its HCl salt after treatment with aq. NaOH and Et2O extr., in MeOH at room temp. to give 2.6 g I (R = H, R1R2 = (CH2)5, R3 = Me, X = iodo, Z = O), which had anticholinergic activity with ED50 of 1.6 + 10-8 g/mL in guinea pigs. Similarly prepd. were 17 addnl. I and their biol. activity given.</p>				
ST	<p>quaternary arom ether antispasmodic; anticholinergic quaternary aryloxypropylammonium halide; antiinflammatant quaternary aryloxypropylammonium halide; analgesic quaternary aryloxypropylammonium halide; aryloxypropylammonium halide antispasmodic anticholinergic; aryloxypropylpiperidinium halide antispasmodic anticholinergic</p>				
IT	<p>Analgesics Inflammation inhibitors Muscle relaxants and Spasmolytics Parasympatholytics (aryloxypropylammonium halides)</p>				
IT	42064-71-1P	42064-72-2P	42064-73-3P	42064-74-4P	42064-76-6P
	42064-79-9P	42064-85-7P	42796-63-4P	42796-67-8P	
	42968-33-0P	51074-51-2P	51543-53-4P	62663-36-9P	
	<p>RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and anticholinergic activity of)</p>				
IT	42063-78-9P	42796-62-3P	42796-71-4P	43213-28-1P	
	51543-52-3P	62663-50-7P			
	<p>RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)</p>				
IT	42064-87-9P	42064-89-1P			
	<p>RL: SPN (Synthetic preparation); PREP (Preparation) (prepn., antispasmodic and analgesic activity of)</p>				
IT	42796-29-2	62663-37-0	62663-38-1	62663-39-2	62663-40-5
	62663-41-6	62663-42-7	62663-43-8	62663-44-9	
	62663-45-0	62663-46-1	62663-47-2	62663-48-3	62663-49-4
	<p>RL: RCT (Reactant) (quaternization of)</p>				
IT	42796-63-4P	62663-36-9P			
	<p>RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and anticholinergic activity of)</p>				
RN	42796-63-4	CAPLUS			
CN	<p>Benzenepropanaminium, N,N-diethyl-N-methyl-γ-phenoxy-, iodide (9CI) (CA INDEX NAME)</p>				



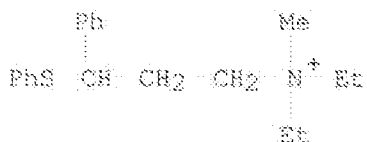
RN 62663-36-9 CAPLUS
 CN Benzenepropanaminium, N,N-diethyl-N-methyl-γ-[(4-methylphenyl)thio]-, iodide (9CI) (CA INDEX NAME)



IT 42796-62-3P 51543-52-3P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (prepn. of)
 RN 42796-62-3 CAPLUS
 CN Benzenepropanaminium, N,N,N-trimethyl-γ-phenoxy-, iodide (9CI) (CA INDEX NAME)



RN 51543-52-3 CAPLUS
 CN Benzenepropanaminium, N,N-diethyl-N-methyl-γ-(phenylthio)-, iodide (9CI) (CA INDEX NAME)



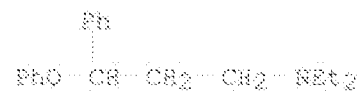
IT 62663-42-7 62663-43-8

RL: RCT (Reactant)

(quaternization of)

RN 62663-42-7 CAPLOS

CN Benzenepropanamine, N,N-diethyl-γ-phenoxy- (9CI) (CA INDEX NAME)



RN 62663-43-8 CAPLOS

CN Benzenepropanamine, N,N-diethyl-γ-[(4-methylphenyl)thio]- (9CI) (CA INDEX NAME)

